Food and Drug Administration, HHS

the premarket notification procedures in subpart E of part 807 of this chapter subject to §880.9.

[45 FR 69682-69737, Oct. 21, 1980, as amended at 63 FR 59229, Nov. 3, 1998]

§ 880.6785 Manual patient transfer device.

- (a) *Identification*. A manual patient transfer device is a device consisting of a wheeled stretcher and a mechanism on which a patient can be placed so that the patient can be transferred with minimal disturbance in a horizontal position to the stretcher.
- (b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter, subject to the limitations in §880.9. The device is also exempt from the current good manufacturing practice regulations in part 820 of this chapter, with the exception of §820.180, with respect to general requirements concerning records, and §820.198, with respect to complaint files.

 $[45~{\rm FR}~69682{-}69737,~{\rm Oct.}~21,~1980,~{\rm as}~{\rm amended}$ at $66~{\rm FR}~38807,~{\rm July}~25,~2001]$

§880.6800 Washers for body waste receptacles.

- (a) Identification. A washer for body waste receptacles is a device intended for medical purposes that is used to clean and sanitize a body waste receptacle, such as a bedpan. The device consists of a wall-mounted plumbing fixture with a door through which a body waste receptacle is inserted. When the door is closed the body waste receptacle is cleaned by hot water, steam, or germicide.
- (b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter, subject to the limitations in §880.9. The device also is exempt from the current good manufacturing practice regulations in part 820 of this chapter, with the exception of §820.180, with respect to general requirements concerning records, and §820.198, with respect to complaint files.

 $[45\ {\rm FR}\ 69682\text{--}69737,\ {\rm Oct.}\ 21,\ 1980,\ {\rm as}\ {\rm amended}$ at $66\ {\rm FR}\ 38807,\ {\rm July}\ 25,\ 2001]$

§880.6820 Medical disposable scissors.

- (a) *Identification*. Medical disposable scissors are disposable type general cutting devices intended for medical purposes. This generic type of device does not include surgical scissors.
- (b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter, subject to the limitations in §880.9.

[45 FR 69682-69737, Oct. 21, 1980, as amended at 66 FR 38807, July 25, 2001]

§880.6850 Sterilization wrap.

- (a) Identification. A sterilization wrap (pack, sterilization wrapper, bag, or accessories, is a device intended to be used to enclose another medical device that is to be sterilized by a health care provider. It is intended to allow sterilization of the enclosed medical device and also to maintain sterility of the enclosed device until used.
- (b) Classification. Class II (performance standards).

§880.6860 Ethylene oxide gas sterilizer.

- (a) *Identification*. An ethylene gas sterilizer is a nonportable device intended for use by a health care provider that uses ethylene oxide (ETO) to sterilize medical products.
- (b) Classification. Class II (performance standards).

§ 880.6870 Dry-heat sterilizer.

- (a) *Identification*. A dry-heat sterilizer is a device that is intended for use by a health care provider to sterilize medical products by means of dry heat.
- (b) Classification. Class II (performance standards).

§880.6880 Steam sterilizer.

- (a) *Identification*. A steam sterilizer (autoclave) is a device that is intended for use by a health care provider to sterilize medical products by means of pressurized steam.
- (b) Classification. Class II (performance standards).

§880.6885 Liquid chemical sterilants/ high level disinfectants.

(a) *Identification*. A liquid chemical sterilant/high level disinfectant is a